VERSION SHOWING MARKED CHANGES

IN THE CLAIMS:

- 1. Canceled.
- 2. Canceled.

(Amended) A compound comprising a metal complexed with a chelating group attached to a gastrin releasing peptide (GRP) receptor agonist, the gastrin releasing peptide receptor agonist including a bombesin agonist binding moiety, said compound having a structure of the formula X-Y-B wherein X is a metal chelating group, Y is a spacer group or covalent bond and B is a gastrin releasing peptide receptor agonist which includes a bombesin agonist binding moiety. The compound of claim 2 wherein and Y is selected from the group consisting of at least one amino acid residue, a hydrocarbon chain and a combination thereof.

(Amended) The compound of claim 2 2 wherein X is selected from the group consisting of DOTA, DTPA, S4, N3S, N2S2, NS3 and derivatives thereof.

(Original) The compound of claim A wherein Y is selected from the group consisting of at least one amino acid residue, a hydrocarbon chain and a combination thereof and B is selected from the group consisting of BBN(7-14) and BBN(8-14).

(Original) The compound of claim A wherein X is DOTA or a derivative thereof.

(Original) The compound of claim wherein Y is selected is selected from the group consisting of at least one amino acid residue, a hydrocarbon chain and a combination thereof and B is selected from the group consisting of BBN(7-14) and BBN(8-14).

(Original) The compound of claim wherein Y is a combination of L-glutamine and a hydrocarbon chain.

(Original) The compound of claim & wherein Y is a combination of L-glutamine and a C1 to C10 hydrocarbon chain.

(Original) The compound of claim wherein Y is selected from the group consisting of glycine, β-alanine, gamma-aminobutanoic acid, 5-aminovaleric acid (5-





Ava), 6-aminohexanoic acid, 7-aminoheptanoic acid, 8-aminooctanoic acid (8-Aoc), 9-aminononanoic acid, 10-aminodecanoic acid and 11-aminoundecanoic acid (11-Aun).

(Original) The compound of claim 4 wherein X is N3S or a derivative thereof.

(Original) The compound of claim wherein Y is selected from the group consisting of at least one amino acid residue, a hydrocarbon chain and a combination thereof and B is selected from the group consisting of BBN(7-14) and BBN(8-14).

(Original) The compound of claim 12 wherein Y is gly-ser-gly.

114. Canceled.

structure of the formula X-Y-B wherein X is a metal chelating group, Y is a spacer group or covalent bond and B is a gastrin releasing peptide (GRP) receptor agonist, the GRP receptor agonist including a bombesin agonist moiety The complex of claim 14 wherein and the metal is selected from the group consisting of transition metals, lanthanides, auger-electron emitting isotopes, and α -, β - or γ -emitting isotopes.

(Amended) The complex of claim 14 15 wherein the metal is selected from the group consisting of: 105Rh-, 99mTc-, 186/188Re-, 153Sm-, 166Ho-, 111In-, 90Y-, 177Lu-, 149Pm-, 166Dy-, 175Yb-, 199Au- and 117mSn-.

(Original) The complex of claim wherein X is selected from the group consisting of DOTA, DTPA, S4, N3S, N2S2, NS3 and derivatives thereof.

(Original) The complex of claim wherein Y is selected from the group consisting of at least one amino acid residue, a hydrocarbon chain and a combination thereof and B is selected from the group consisting of BBN(7-14) and BBN(8-14).

(Original) The complex of claim wherein X is DOTA or a derivative thereof.

(Original) The complex of claim 19 wherein Y is selected is selected from the group consisting of at least one amino acid residue, a hydrocarbon chain and a combination thereof and B is selected from the group consisting of BBN(7-14) and BBN(8-14).

(Original) The complex of claim 20 wherein Y is a combination of L-glutamine and a hydrocarbon chain.



(Original) The complex of claim 21 wherein Y is a combination of L-glutamine and a C1 to C10 hydrocarbon chain.

23. (Original) The complex of claim 22 wherein Y is selected from the group consisting of glycine, β-alanine, gamma-aminobutanoic acid, 5-aminovaleric acid (5-Ava), 6-aminohexanoic acid, 7-aminoheptanoic acid, 8-aminooctanoic acid (8-Aoc), 9-aminononanoic acid, 10-aminodecanoic acid and 11-aminoundecanoic acid (11-Aun).

(Original) The complex of claim 23 wherein Y is 8-aminooctanoic acid.

(Original) The complex of claim 23 consisting of 90Y-DOTA-8-Aoc-BBN(7-14)NH2.

13,26. (Original) The complex of claim 23 consisting of 111In-DOTA-8-Aoc-BBN(7-14) NH2.

(Original) The complex of claim 25 consisting of 177Lu-DOTA-8-Aoc-BBN(7-14) NH2.

BBN(7-14) NH2.

(Original) The complex of claim 28 consisting of 90Y-DOTA-5-Ava-BBN(7-14)NH2.

30. (Original) The complex of claim 23 consisting of 111In-DOTA-5-Ava-BBN(7-14) NH2.

2831. (Original) The complex of claim 23 consisting of 177Lu-DOTA-5-Ava-BBN(7-14) NH2.

9732. (Original) The complex of claim 23 consisting of 149Pm-DOTA-5-Ava-BBN(7-14) NH2.

(Original) The complex of claim of wherein X is N3S or a derivative thereof.

(Original) The complex of claim 33 wherein Y is selected from the group consisting of at least one amino acid residue, a hydrocarbon chain and a combination thereof and B is selected from the group consisting of BBN(7-14) and BBN(8-14).

7 35. (Original) The complex of claim 34 wherein Y is gly-ser-gly.

39.36. (Original) The complex of claim 34 consisting of 99mTc-N3S-gly-ser-gly-BBN(7-14)NH2.

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37. Canceled.

A method of treating patients using radioisotope therapy by administering an effective amount of a pharmaceutical comprising a metal complex with a chelating group with a GRP receptor agonist, the GRP receptor agonist including a bombesin agonist moiety. The method according to claim 37, wherein said method includes administering an effective amount of a the complex comprising a metal and a compound having a structure of the formula X-Y-B wherein X is a metal chelating group, Y is a spacer group or covalent bond and B is a gastrin releasing peptide receptor agonist which includes a bombesin agonist binding moiety.

(Original) The method of claim 38 wherein the metal is selected from the group consisting of transition metals, lanthanides, auger-electron emitting isotopes, and α -, β - or γ -emitting isotopes.

(Original) The method of claim 38 wherein the metal is selected from the group consisting of: 105Rh-, 99mTc-, 186/188Re-, 153Sm-, 166Ho-, 111In-, 90Y-, 177Lu-, 149Pm-, 166Dy-, 175Yb-, 199Au- and 1,17mSn-.

31°41. (Original) The method of claim wherein X is selected from the group consisting of DOTA, DTPA, S4, N3S, N2S2, NS3 and derivatives thereof.

38 AZ. (Original) The method of claim 4 wherein X is DOTA or a derivative thereof.

39 48. (Original) The method of claim 22 wherein Y is selected from the group consisting of at least one amino acid residue, a hydrocarbon chain and a combination thereof and B is selected from the group consisting of BBN(7-14) and BBN(8-14).

(Original) The method of claim 32 wherein Y is a combination of L-glutamine and a hydrocarbon chain.

(Original) The method of claim 44 wherein Y is selected from the group consisting of glycine, β-alanine, gamma-aminobutanoic acid, 5-aminovaleric acid (5-Ava), 6-aminohexanoic acid, 7-aminoheptanoic acid, 8-aminooctanoic acid (8-Aoc), 9-aminononanoic acid, 10-aminodecanoic acid and 11-aminoundecanoic acid (11-Aun).

(Original) A method of imaging a patient by administering to a subject a diagnostically effective amount of a compound as set forth in claim.

(Original) The method of claim 46, wherein said method includes administering an effective amount of a complex comprising a metal and a compound having a structure of the formula X-Y-B wherein X is a metal chelating group, Y is a spacer group or covalent bond and B is a gastrin releasing peptide receptor agonist which includes a bombesin agonist binding mojety.

(Original) The method of claim $\frac{2}{3}$ wherein the metal is selected from the group consisting of transition metals, lanthanides, auger-electron emitting isotopes, and α -, β - or y-emitting isotopes.

(Original) The method of claim As wherein X is selected from the group consisting of DOTA, DTPA, S4, N3S, N2S2, NS3 and derivatives thereof.

the 50. (Original) The method of claim claim wherein X is N3S or a derivative thereof.

(Original) The method of claim 50 wherein Y is selected is selected from the group consisting of at least one amino acid residue, a hydrocarbon chain and a combination thereof and B is selected from the group consisting of BBN(7-14) and BBN(8-14).

(Original) The method of claim 54 wherein Y is gly-ser-gly.

(Previously Amended) A method of forming a therapeutic or diagnostic compound comprising the step of reacting a metal complexed with a chelating group with a GRP receptor agonist the receptor agonist including a bombesin agonist moiety.

(Original) The method of claim 53, wherein said method includes reacting a metal with a compound having a structure of the formula X-Y-B wherein X is a metal chelating group, Y is a spacer group or covalent bond and B is a gastrin releasing peptide receptor agonist which includes a bombesin agonist binding moiety.

955. (Original) The method of claim 54 wherein the metal is selected from the group consisting of transition metals, lanthanides, auger-electron emitting isotopes, and α -, β - or y-emitting isotopes.

5.56. (Original) The method of claim 54 wherein the metal is selected from the group consisting of: 99mTc- and 186/188Re-.

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(Original) The method of claim 36 wherein Y is selected is selected from the group consisting of at least one amino acid residue, a hydrocarbon chain and a combination thereof.

(Original) The method of claim 57 wherein X is selected from the group consisting of DOTA, DTPA, S4, N3S, N2S2, NS3 and derivatives thereof.

(Original) The method of claim 55 wherein B is selected from the group consisting of BBN(7-14) and BBN(8-14).

be 60. (Original) The method of claim 59 wherein X is DOTA or a derivative thereof and Y is selected from the group consisting of glycine, β-alanine, gamma-aminobutanoic acid, 5-aminovaleric acid (5-Ava), 6-aminohexanoic acid, 7-aminohexanoic acid, 8-aminooctanoic acid (8-Aoc), 9-aminononanoic acid, 10-aminodecanoic acid and 11-aminoundecanoic acid (11-Aun).

51 (Original) The method of claim 55 wherein X is N3S or a derivative thereof and Y is gly-ser-gly.



